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THE 10TH SICKLE CELL IN FOCUS 2 - 3 JUNE 2016 CONFERENCE

Natcher Conference Center
National Heart, Lung and Blood Institute (NHLBI)
National Institutes of Health (NIH)
Bethesda, near Washington DC, USA

Co-hosted by the **National Heart, Lung and Blood Institute** and the **South Thames Sickle Cell & Thalassaemia Network**, this two-day, intensive, educational conference highlights and discusses emerging clinical complications and management of sickle cell disease.

The clinical and scientific lectures are aimed at consultants, trainee doctors, healthcare professionals involved in the care of patients with sickle cell disease, plus academic researchers in this field.

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SESSION ONE: SICKLE CELL DISEASE - A GLOBAL HEALTH ISSUE

United States of America:

Kathy Hassell

University of Colorado, USA

Africa:

Dapa Diallo

University of Bamako, Mali

Brazil:

Anna Proietti

Hemominas Foundation, Brazil

Europe:

Beatrice Gulbis

Hôpital Erasme, Belgium

The Caribbean:

Jennifer Knight

University Hospital of the West Indies, Jamaica

India:

Roshan B. Colah

National Institute of Immunohaematology,
India

SESSION TWO: SICKLE CELL DISEASE AND MALARIA - MORE THAN JUST A BALANCING ACT

**Protective role of HbS and other red
blood cell defects on malaria:**

Tom Williams

Imperial College London, UK \ KEMRI-
Wellcome Trust Collaborative Programme,
Kilifi, Kenya

**Adhesion, occlusion, and tissue
perfusion in malaria and sickle cell
disease:**

Hans Ackerman

National Heart, Lung and Blood Institute,
National Institutes of Health, USA

**A microfluidic platform to assess
malaria and sickle stickiness:**

Subra Suresh

Carnegie Mellon University, Pittsburgh, USA

SESSION THREE: VASCULAR INJURY & INFLAMMATION IN SICKLE CELL DISEASE

Inflammasomes in health and disease:

Fayyaz Sutterwala

University of Iowa, USA

**Drivers of inflammation in sickle cell
disease - role of TLR:**

John Belcher

University of Minnesota, USA

**Drivers of inflammation in sickle cell
disease - role of neutrophils:**

Dachuan Zhang

Albert Einstein College of Medicine, USA

SESSION FOUR: UPDATES

**Newborn screening and point-of-
care testing in developing countries:**

Carolyn Hoppe

Children's Hospital & Research Center,
USA

Clinical trials:

Deepa Manwani

The Children's Hospital at Montefiore, USA

Bone marrow transplant:

John Horan

Children's Healthcare of Atlanta, USA

SESSION FIVE: DEBATE
**HSCT IS RECOMMENDED FOR SECONDARY PREVENTION OF CEREBROVASCULAR
INFARCTS IN SICKLE CELL DISEASE**

For:

Michael DeBaun

Vanderbilt University School of Medicine, USA

Against:

Elliott Vichinsky

Children's Hospital & Research Center, Oakland, USA

SESSION ONE: GENETICS & GENOMICS - PRIME TIME IN HEMOGLOBIN DISORDERS

Genetics and genomics in clinical studies –potentials and pitfalls:

Steve Chanock

National Cancer Institute, National Institutes of Health, USA

Impact of genetics and genomics in sickle cell disease:

Swee Lay Thein

National Heart, Lung and Blood Institute, National Institutes of Health, USA

Gene therapy:

John Tisdale

National Heart, Lung and Blood Institute, National Institutes of Health, USA

Application of genome editing in hemoglobin disorders:

Dan Bauer

Children's Hospital Boston, USA

Environmental determinants of disease severity:

David Rees

King's College Hospital NHS Foundation Trust / King's College London, UK

SESSION TWO: UPDATES ON MANAGEMENT - PART ONE

Management of pain and opioid use in patients with sickle cell disease:

Patrick Carroll

Johns Hopkins University School of Medicine, USA

Inflammation and pain in sickle cell disease:

Deepika Darbari

Children's National Medical Center, USA

Causes, complications and management of hypoxia in sickle cell disease:

Roberto Machado

University of Illinois College of Medicine at Chicago, USA

SESSION THREE : UPDATES ON MANAGEMENT - PART TWO

Expanding use of hydroxyurea in sickle cell disease:

Jane Hankins

St Jude Children's Research Hospital, USA

Blood transfusion, alloimmunisation and the role of variant genotyping:

Stella Chou

The Children's Hospital of Philadelphia, USA

VTEs in sickle cell disease:

Kenneth Ataga

University of North Carolina, USA

SESSION FOUR: DEBATE

HIGH MORTALITY IN YOUNG ADULTS IS DUE TO THE BREAKDOWN IN TRANSITION TO ADULT CARE

Yes:

Mariane de Montalembert

Necker-Enfants Malades Hospital, France

No:

Sophie Lanzkron

Johns Hopkins University School of Medicine, Baltimore, USA